



**PRIOR AUTHORIZATION REQUEST FORM**  
**Abemaciclib (Verzenio®)/ Advanced or metastatic breast cancer**

**DECLARATION OF THE INSURED PERSON**

**Section 1: Information about the plan member and the patient**

|                                  |                                |                  |             |
|----------------------------------|--------------------------------|------------------|-------------|
| Name of plan member              | Insurance policy / certificate | Name of employer |             |
| Name of patient                  | Date of birth (YYYY/MM/DD)     | Telephone        |             |
| Address (number and street name) | City/Town                      | Province         | Postal code |

**Section 2: Other prescription drug insurance policies**

|   |                                       |                                  |
|---|---------------------------------------|----------------------------------|
| Do you have other prescription drug insurance?                      | <input type="checkbox"/> Yes          | <input type="checkbox"/> No      |
| If so, please answer the following:                                 |                                       |                                  |
| What type of plan is it?  | <input type="checkbox"/> Private      | <input type="checkbox"/> Public  |
| Have you ever submitted a claim for this drug to the other insurer? | <input type="checkbox"/> Yes          | <input type="checkbox"/> No      |
| What is the status of the claim?                                    | <input type="checkbox"/> Accepted     | <input type="checkbox"/> Refused |
| Did this insurer ask you to complete a prior authorization request? | <input type="checkbox"/> Yes          | <input type="checkbox"/> No      |
| If so, what is the status of the prior authorization request?       | <input type="checkbox"/> Accepted     | <input type="checkbox"/> Refused |
|   | <input type="checkbox"/> Under review |                                  |

***Please enclose acceptance or refusal documents, if applicable***

**Section 3: Authorization to disclose personal information**

I certify that the information in this prior authorization request is complete, accurate and true.

I authorize physicians and other health care professionals, medical, paramedical or clinical institutions, care coordinators, members of SSQ's Preferred Pharmacy Network (outside Quebec only) and any public or parapublic organization, including Régie de l'assurance maladie du Québec, to disclose to SSQ, Life Insurance Company Inc. (SSQ) any of my relevant personal information including and without limitation, any medical information and medical evaluations in connection with the processing of this request. I hereby waive their confidentiality obligation and authorize them to disclose the requested information to SSQ. In addition, I authorize SSQ to disclose to the previously named third parties any of my relevant personal information including and without limitation, any medical information and medical evaluations in connection with the processing of this request.

Photocopies of this document have the same value as the original.

Signature of **patient** (parent/legal guardian) \_\_\_\_\_ Date \_\_\_\_\_

**IMPORTANT:**

All correspondence concerning this form will be sent to the address indicated in the participant's file.

**Send us this duly completed form by mail or by fax to: 1-855-453-3942.**

Telephone: 418-651-2588/1-800-380-2588 – Fax: 1-855-453-3942 Address: 2525 Laurier Blvd, P.O. Box 10500, Quebec City, QC G1V 4H6 /ssq.ca



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**DECLARATION OF THE PRESCRIBER**

| Section 4: Information about the prescriber   |           |               |
|---|-----------|---------------|
| Name of prescriber  | Specialty | License no.   |
| Telephone   | Fax       |               |
| I hereby certify that the information in this request is complete, true and accurate. |           |               |
| _____<br>Signature of <b>prescriber</b>   |           | _____<br>Date |

| Section 5: Drug covered by the authorization |   |          |  |
|--|---|----------|--|
| Name of drug                                 | Pharmaceutical form   | Strength | <b>Dosage</b><br>Dose: _____<br>Frequency of administration: _____ |
| Type of request                              | <input type="checkbox"/> First request<br>Complete section 6  |          |  |
|  | <input type="checkbox"/> Continuation of treatment<br>Complete section 7<br>Also complete section 6 if this is the first authorization requested from SSQ |          |  |



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**IMPORTANT:**

**Please do not provide any genetic test results**

**Section 6: Clinical information (first request)**

**Diagnosis**

☐ Advanced or metastatic breast cancer in compliance with Health Canada Indication

For informational purposes only:

VERZENIO® (abemaciclib) is indicated by Health Canada for the treatment of hormone receptor (HR +)-positive, human epidermal growth factor receptor 2 (HER2-)-negative advanced or metastatic breast cancer:

- in combination with an aromatase inhibitor in postmenopausal women as initial endocrine-based therapy.
- in combination with fulvestrant in women with disease progression following endocrine therapy. Pre- or perimenopausal women must also be treated with a gonadotropin-releasing hormone (GnRH) agonist.
- as a single agent in women with disease progression following endocrine therapy and at least 2 prior chemotherapy regimens. At least one chemotherapy regimen should have been administered in the metastatic setting, and at least one should have contained a taxane.

☐ Other, specify: \_\_\_\_\_

**Complete the following information**

☐ Post-menopausal

☐ Pre/Perimenopausal

Actual value of the ECOG performance status

☐ 0

☐ 1

☐ 2

☐ 3

☐ 4

**Administration of Verzenio®**

Administered as first-line metastatic treatment?

☐ Yes

☐ No

☐ In association with an aromatase inhibitor. Specify: \_\_\_\_\_

☐ In association with Fulvestrant

☐ In monotherapy

☐ Other, specify: \_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_



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| Section 6: Clinical information (first request) (cont'd) |   |                        |
|--|---|------------------------|
| Summary of previous trials or contraindications          |   |                        |
| Drug or other medical treatment                          | Reason for discontinuation  | Duration of treatment  |
| Name: _____<br>Dosage: _____                             | <input type="checkbox"/> Ineffectiveness<br><input type="checkbox"/> Intolerance<br><input type="checkbox"/> Contraindication<br><input type="checkbox"/> Other, specify: _____ | From _____<br>To _____ |
| Name: _____<br>Dosage: _____                             | <input type="checkbox"/> Ineffectiveness<br><input type="checkbox"/> Intolerance<br><input type="checkbox"/> Contraindication<br><input type="checkbox"/> Other, specify: _____ | From _____<br>To _____ |
| Name: _____<br>Dosage: _____                             | <input type="checkbox"/> Ineffectiveness<br><input type="checkbox"/> Intolerance<br><input type="checkbox"/> Contraindication<br><input type="checkbox"/> Other, specify: _____ | From _____<br>To _____ |
| Name: _____<br>Dosage: _____                             | <input type="checkbox"/> Ineffectiveness<br><input type="checkbox"/> Intolerance<br><input type="checkbox"/> Contraindication<br><input type="checkbox"/> Other, specify: _____ | From _____<br>To _____ |
| Name: _____<br>Dosage: _____                             | <input type="checkbox"/> Ineffectiveness<br><input type="checkbox"/> Intolerance<br><input type="checkbox"/> Contraindication<br><input type="checkbox"/> Other, specify: _____ | From _____<br>To _____ |

## Section 7: Clinical information (continuation of treatment)

### Information necessary to evaluate the response to treatment

The drug covered by the present authorization request was first taken on (YYYY-MM-DD): \_\_\_\_\_

### Positive clinical effects observed

Date treatment began (YYYY-MM-DD): \_\_\_\_\_

- ☐ Absence of disease progression
- ☐ Other, specify: \_\_\_\_\_

## Section 8: Additional information

[illegible]