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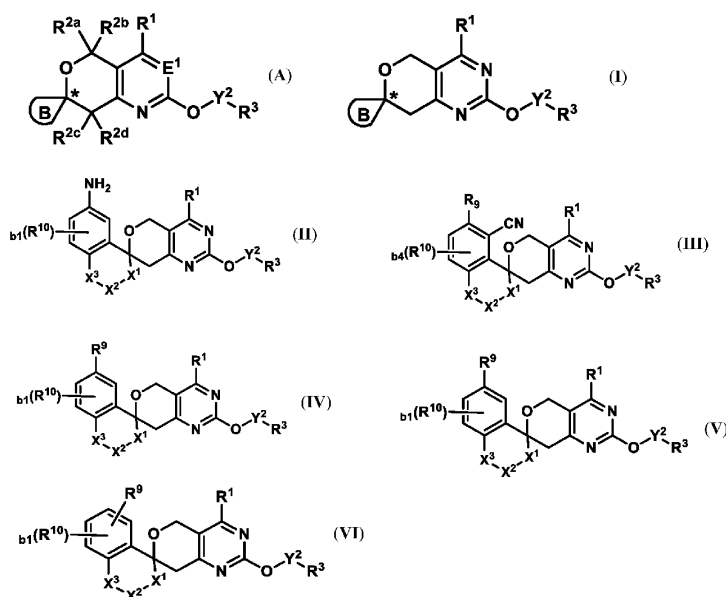
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- (71) Applicant: **TREELINE BIOSCIENCES, INC.** [US/US];
677 Washington Blvd., Suite 525, Stamford, Connecticut
06901 (US).
- (72) Inventors: **CABRAL, Shawn**; 500 Arsenal Street, 2nd
Floor, Watertown, Massachusetts 02472 (US). **MASCIT-
TI, Vincent**; 500 Arsenal Street, 2nd Floor, Watertown,
Massachusetts 02472 (US). **DECULTOT, Ludovic Jacky
Gilbert**; 500 Arsenal Street, 2nd Floor, Watertown, Mass-
achusetts 02472 (US). **HEROLD, Jens-Martin**; 500 Ar-
senal Street, 2nd Floor, Watertown, Massachusetts 02472
(US). **HESP, Kevin D.**; 500 Arsenal Street, 2nd Floor, Wa-
tertown, Massachusetts 02472 (US). **LAJINESS, James
Paul**; 11180 Roselle Street, Suite A, San Diego, Califor-
nia 92121 (US). **SEIM, Alexander E.**; 500 Arsenal Street,
2nd Floor, Watertown, Massachusetts 02472 (US). **TO-
FOLEANU, Florentina**; 500 Arsenal Street, 2nd Floor,

(54) Title: SPIROCYCLIC DIHYDROPYRANOPYRIMIDINE KRAS INHIBITORS



(57) Abstract: This disclosure provides compounds of Formula (A) (e.g., Formula (I) (e.g., Formula (I-a1), (I-a2), (I-a3), (I-a4), (I-a5), (I-b1), (I-b2), (I-b3), (I-b4), (I-b5), or (I-c1)), Formula (II) (e.g., Formula (II-a), (II-b), (II-a1), (II-b1), (II-a2), or (II-b2)), Formula (III) (e.g., Formula (III-1) or (III-2)), Formula (IV) (e.g., Formula (IV-a), (IV-b), (IV-c), (IV-a1), (IV-b1), (IV-a2), or (IV-b2)), or Formula (V) (e.g., Formula (V-a) or (V-b), (V-a1), (V-c), (V-d), (V-b1), (V-a2), or (V-b2)), or Formula (VI) (e.g., Formula (VI-a), (VI-b), (VI-c), (VI-d), or (VI-e))), or pharmaceutically acceptable salts thereof, that inhibit a KRas protein. These compounds are useful, for example, for treating a disease, disorder, or condition in which increased and/or sustained KRas activation (e.g., KRas activation associated with a mutant KRas protein), contributes to the pathology and/or symptoms and/or progression of the disease, disorder, or condition. This disclosure also provides compositions containing the compounds provided herein, or pharmaceutically acceptable salts thereof.

[Continued on next page]

Watertown, Massachusetts 02472 (US). **TSAI, Andy**; 500 Arsenal Street, 2nd Floor, Watertown, Massachusetts 02472 (US).

(74) **Agent: KENDALL, John T.** et al.; Fish & Richardson P.C., P.O. Box 1022, Minneapolis, Minnesota 55440-1022 (US).

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